IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

Application of James T. English et al.

Art Unit 1639

Serial No.: 09/829,549 Filed: April 10, 2001 Confirmation No. 8198

For PHAGE DISPLAY SELECTION OF ANTI FUNGAL PEPTIDES

Examiner: Teresa D. Wessendorf

Commissioner for Patents P.O. Box 1450 Alexandria VA 22313-1450

November 10, 2004

AMENDMENT D

Sir:

In response to the Office Action mailed June 10, 2004, and reinstated September 13, 2004, please enter the following amendments and consider the following remarks set forth in this Amendment D and Response After Request For Continued Examination.

Amendments to the Claims are reflected in the listing of claims which begins on page 2 of this paper.

Remarks begin on page 7 of this paper.

Conclusion begins on page 13 of this paper.

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AMENDMENTS TO THE CLAIMS:

This listing of claims will replace all prior versions, and listings, of claims in the application:

LISTING OF CLAIMS:

- 1. (previously presented) A method for identification of non-immunoglobulin peptides having an affinity for the surface of a fungus comprising:
 - (a) constructing a library of peptides by,
 - (i) preparing random oligonucleotides;
- (ii) inserting said oligonucleotides into a vector that expresses peptides encoded by said random oligonucleotides on its surface and is capable of transfecting a host cell;
- (iii) transfecting a host cell with said vector to amplify said vector in an infectious form to create a library of peptides on the surface of said vector;
- (b) contacting said vector expressing said peptide library with a target fungus and removing unbound vector;
 - (c) eluting bound vector from said fungus;
 - (d) amplifying said bound vector;
 - (e) sequencing the oligonucleotides contained in said eluted vector;
- (f) deducing the amino acid sequence of peptides encoded by said oligonucleotides contained in said eluted vector; and
- (g) selecting the non-immunoglobulin peptides for which the amino acid sequence has been deduced.
- 2. (currently amended) The method of <u>any one of claims</u> 1, 48, or 49 further comprising repeating steps (b) through (d) at least once.
- 3. (currently amended) The method of <u>any one of claims</u> 1, 48, or 49, wherein said vector is a fusion phage vector.

- 4. (currently amended) The method of <u>any one of claims</u> 1, 48, or 49, wherein said vector is a fusion phage vector selected from the group consisting of type 8, type 88, type 8+8, type 3, type 3+3, type 6, type 66, type 6+6, phage T7 and phage 8.
- 5. (currently amended) The method of <u>any one of claims</u> 1 <u>or 48</u>, wherein the sequence of said random oligonucleotide is GCA GNN (NNN)7 or SEQ ID NO: 1.
- 6. (currently amended) The method of <u>any one of claims</u> 1, 48, or 49, wherein said peptide is expressed as part of a coat protein of said vector.
- 7. (original) The method of claim 6, wherein said coat protein is a plll or a pVIII coat protein.
- 8. (currently amended) The method of <u>any one of claims</u> 1, 48, or 49, further comprising determining the binding affinity of said peptides to said target fungus.
- 9. (currently amended) The method of <u>any one of claims</u> 1 <u>or 48</u>, wherein each of said peptides are of the same length, the length being 6 to 15 amino acids.

10-31. (canceled)

- 32. (currently amended) The method of <u>any one of claims</u> 1 <u>or 49</u> wherein the target fungus is a plant pathogenic fungus.
- 33. (currently amended) The method of <u>any one of claims</u> 1 <u>or 49</u> wherein the target fungus is a member of genus *Phytophthora*.
- 34. (currently amended) The method of <u>any one of claims</u> 1<u>or 49</u> wherein the target fungus is selected from the group consisting of *Phytophthora sojae*, *Phytophthora*

capsici, Phytophthora cactorum, Phytophthora palmivora, Phytophthora cinnamomi, Phytophthora infestans, and Phytophthora parasitica.

- 35. (currently amended) The method of <u>any one of claims</u> 1 <u>or 49</u> wherein the target fungus is selected from the group consisting of *Phytophthora sojae*, *Phytophthora capsici*, *Phytophthora palmivora*, *Phytophthora cinnamomi*, and *Phytophthora parasitica*.
- 36. (currently amended) The method of <u>any one of claims</u> 1 or 49 wherein the target fungus is *Phytophthora sojae* or *Phytophthora capsici*.
- 37. (currently amended) The method of <u>any one of claims</u> 1, 48, or 49 wherein the vector expressing the peptide library is contacted with the target fungus at different life stages of the target fungus.
- 38. (currently amended) The method of <u>any one of claims</u> 1, 48, or 49 wherein the vector expressing the peptide library is contacted with the target fungus at oospore life stage or chlamydospore life stage.
- 39. (currently amended) The method of <u>any one of claims</u> 1, 48, or 49 wherein the vector expressing the peptide library is contacted with the target fungus at zoospore life stage.
- 40. (currently amended) The method of <u>any one of claims</u> 1, 48, or 49 wherein the vector expressing the peptide library is contacted with the target fungus at germling life stage.
- 41. (currently amended) The method of <u>any one of claims</u> 1 <u>or 48</u> wherein each of said peptides are of a same length, the length being 8 amino acids.

- 42. (currently amended) The method of <u>any one of claims</u> 1 <u>or 48</u> wherein the peptide library is an f8-1 peptide library.
- 43. (currently amended) The method of <u>any one of claims 1 or 48</u> wherein each of said peptides are of a same length, the length being 15 amino acids.
- 44. (currently amended) The method of <u>any one of claims</u> 1 <u>or 48</u> wherein the peptide library is an f88-4 peptide library.
- 45. (currently amended) The method of <u>any one of claims</u> 1, 48, or 49, further comprising repeating steps (b) through (d) at least twice.
- 46. (currently amended) The method of <u>any one of claims</u> 1, 48, or 49, further comprising repeating steps (b) through (d) at least three times.
- 47. (currently amended) The method of <u>any one of claims</u> 1, 48, or 49 wherein the bound vector is amplified in an *E. coli*.

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(new) A method for identification of non-immunoglobulin peptides having an affinity for the surface of a fungus comprising:

- (a) constructing a library of peptides by,
 - (i) preparing random oligonucleotides;
- (ii) inserting said oligonucleotides into a vector that expresses peptides encoded by said random oligonucleotides on its surface and is capable of transfecting a host cell;
- (iii) transfecting a host cell with said vector to amplify said vector in an infectious form to create a library of peptides on the surface of said vector;
- (b) contacting said vector expressing said peptide library with a target fungus and removing unbound vector, wherein the target fungus is selected from the group consisting of *Phytophthora sojae*, *Phytophthora capsici*, *Phytophthora palmivora*, *Phytophthora cinnamomi*, and *Phytophthora parasitica*;

- (c) eluting bound vector from said fungus;
- (d) amplifying said bound vector;
- (e) sequencing the oligonucleotides contained in said eluted vector;
- (f) deducing the amino acid sequence of peptides encoded by said oligonucleotides contained in said eluted vector; and
- (g) selecting the non-immunoglobulin peptides for which the amino acid sequence has been deduced.

75
Ast. (new) A method for identification of non-immunoglobulin peptides having an affinity for the surface of a fungus comprising:

- (a) constructing a library of peptides by,
 - (i) preparing random oligonucleotides;
- (ii) inserting said oligonucleotides into a vector that expresses peptides encoded by said random oligonucleotides on its surface and is capable of transfecting a host cell;
- (iii) transfecting a host cell with said vector to amplify said vector in an infectious form to create a library of peptides on the surface of said vector;

wherein the library of peptides is (1) an f8-1 peptide library, wherein each peptide of the f8-1 peptide library has a length of 8 amino acids or (2) an f88-4 peptide library, wherein each peptide of the f88-4 peptide library has a length of 15 amino acids;

- (b) contacting said vector expressing said peptide library with a target fungus and removing unbound vector;
 - (c) eluting bound vector from said fungus;
 - (d) amplifying said bound vector;
 - (e) sequencing the oligonucleotides contained in said eluted vector;
- (f) deducing the amino acid sequence of peptides encoded by said oligonucleotides contained in said eluted vector; and
- (g) selecting the non-immunoglobulin peptides for which the amino acid sequence has been deduced.